

Experimental Determination and System Level Analysis of Essential Genes in *Escherichia coli* MG1655

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Defining the gene products that play an essential role in an organism's functional repertoire is vital to understanding the system level organization of living cells. We used a genetic footprinting technique for a genome-wide assessment of genes required for robust aerobic growth of *Escherichia coli* in rich media. We identified 620 genes as essential and 3,126 genes as dispensable for growth under these conditions. Functional context analysis of these data allows individual functional assignments to be refined. Evolutionary context analysis demonstrates a significant tendency of essential *E. coli* genes to be preserved throughout the bacterial kingdom. Projection of these data over metabolic subsystems reveals topologic modules with essential and evolutionarily preserved enzymes with reduced capacity for error tolerance.

Sequencing and comparative analysis of multiple diverse genomes is revolutionizing contemporary biology by providing a framework for interpreting and predicting the physiologic properties of an organism. A variety of emerging postgenomic techniques such as genome-wide expression profiling and monitoring of macromolecular complex formation can reveal the detailed molecular compositions of cells. New computational approaches to exploring the inherent organization of cellular networks, the mode and dynamics of interactions among cellular constituents, are in early stages of development (14, 22, 23). These techniques allow us to begin unraveling a major paradigm of cellular biology: how biological properties arise from the large number of components making up an individual cell.

Defining which gene products play an essential role and under what conditions is vital to understanding the complexity of living organisms. Although methods to rapidly and systematically determine genome-wide gene essentiality are less advanced than other functional genomic techniques, a number of essentiality surveys involving different species have been reported. Many experimental approaches have been used to produce such data, including individual knockouts in *Saccharomyces cerevisiae* (10, 38), *Caenorhabditis elegans* (21), and recently *B. subtilis* (22a), RNA interference in *C. elegans* (20), and whole-genome transposon mutagenesis studies with several microorganisms. In the latter group, complete or extensive lists of essential and dispensable genes are available for *Myc-*

plasma pneumoniae and *Mycoplasma genitalium* (15), *Mycobacterium tuberculosis* (31), *Haemophilus influenzae* (1), and *S. cerevisiae* (30). However, as of yet relatively little effort has been committed to a system level interpretation of these data in terms of cellular function or evolutionary relationships with other organisms (19).

Escherichia coli has historically been the focus of intense biochemical, genetic, and physiologic scrutiny, but genomic essentiality data for this organism have remained incomplete. Systematic efforts to compile genome-wide collections of *E. coli* deletion mutants are under way. Two groups have reported Tn10 transposon-based genetic footprinting projects with *E. coli*, but essentiality data were revealed only for a limited set of genes (3, 13). Currently, the Profiling of *E. coli* Chromosome database (available at <http://www.shigen.nig.ac.jp/ecoli/pec>) is the most complete list of essential and dispensable genes in *E. coli*. This list is not based on direct experimental evidence but is derived from systematic review of the experimental literature. Although this compilation is of great value, the wide variety of strains, conditions, and types of mutations used in individual studies significantly complicates interpretation.

Here we report a genome-wide, comprehensive experimental assessment of the *E. coli* MG1655 genes necessary for robust aerobic growth in a rich, tryptone-based medium. Of the 4,291 protein-encoding genes in *E. coli*, we assessed the essentiality of 3,746 genes (~87% of the total). Individual assessments were projected onto a whole-cell functional reconstruction model including both metabolic and nonmetabolic systems. Distribution of conditionally essential and dispensable *E. coli* genes within functional systems was analyzed with respect to the occurrence of putative orthologs across a broad range of diverse bacterial genomes. This analysis demonstrates a significant tendency of experimentally identified essential *E. coli* genes to be evolutionarily preserved throughout the bac-

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terial kingdom, especially a subset of genes representing key cellular processes such as DNA replication and protein synthesis. Finally, we analyzed the conditional essentiality of metabolic enzymes from the perspective of cellular system level organization, demonstrating enrichment with those enzymes that catalyze reactions within evolutionarily conserved topologic modules in the complex metabolic web of *E. coli*.

MATERIALS AND METHODS

Transposon mutagenesis. *E. coli* strain MG1655 ($F^- \lambda^- \text{ } \textit{ilvG rfb-50 rph-1}$) (16) was used throughout this work. Genetic footprinting with the use of the plasmid pMOD<MCS> containing the artificial transposon EZ::TN<KAN-2> (Epicentre Technologies, Madison, Wis.) and identification of chromosomal insertion sites were previously described (9) and are detailed in the supplementary data (supplementary data for this paper are available at http://www.integratedgenomics.com/online_material/gerdes and on the University of Notre Dame and Northwestern University websites [<http://www.umsl.edu/~balazsi/JBact2003/> and <http://www.oltvailab.northwestern.edu/Pubs/JBact2003/>]). Cells were grown in an enriched Luria-Bertani (LB) medium composed of 10 g of tryptone/liter, 5 g of yeast extract/liter, 50 mM NaCl, 9.5 mM NH_4Cl , 0.528 mM MgCl_2 , 0.276 mM K_2SO_4 , 0.01 mM FeSO_4 , 5×10^{-4} mM CaCl_2 , and 1.32 mM K_2HPO_4 . The growth medium also included the following micronutrients: 3×10^{-6} mM $(\text{NH}_4)_6(\text{MoO}_7)_{24}$, 4×10^{-4} mM H_3BO_3 , 3×10^{-5} mM CoCl_2 , 10^{-5} mM CuSO_4 , 8×10^{-5} mM MnCl_2 , and 10^{-5} mM ZnSO_4 . The following vitamins were added (concentrations are in milligrams per liter): biotin, 0.12; riboflavin, 0.8; pantothenic acid, 10.8; niacinamide, 12.0; pyridoxine, 2.8; thiamine, 4.0; liponic acid, 2.0; folic acid, 0.08; and *p*-aminobenzoic acid, 1.37. Kanamycin was added to 10 $\mu\text{g}/\text{ml}$.

As with any high-throughput technique, genetic footprinting is subject to a certain degree of experimental and analytical error. A variety of validation techniques indicate the overall error rate of our assignments to be well within 10% (9). The actual experimental detection and insert mapping error rate is much lower (within 1 to 2%). The major source of ambiguity is associated with data interpretation (see below). In the supplementary data, we include the insert distribution within each open reading frame (ORF) (raw data, including insert distribution within intergenic regions, are available upon request).

Statistical analyses of transposon insertion frequency. Essential and ambiguous ORFs introduce a bias into the density of transposon insertions due to the fact that they "lose" the insertions incorporated within them during selective outgrowth. There were also unmapped genomic regions where transposon insertions could not be detected. To reconstruct insert distribution prior to selective outgrowth, and to account for the contribution of unmapped regions, we removed from the *E. coli* chromosomal map every ORF with a function asserted to be essential, ambiguous, or not determined, as well as the regions not covered by the mapping process, and joined together the rest of the chromosome. We analyzed the original and corrected insertion location data assuming that the insertions appear as a result of a Poisson process with an overall rate r of 3.218/kb. Based on this hypothesis, the probability to find M insertions within a DNA region of length L is given by

$$P_M(L) = \frac{(rL)^M}{M!} \cdot e^{-rL}$$

The P values corresponding to this hypothesis for the corrected data were calculated to estimate the statistical significance of the deviations from a Poisson process, for a threshold of P of 10^{-5} (see Fig. 1).

If the insertion locations are approximated by a Poisson process, the statistical reliability of essentiality calls depends on two factors: the overall insertion density r in the region where the ORF is located and the length L of the ORF. More specifically, the probability that an ORF is missed by chance is given as follows: $P_0(L) = e^{-rL}$, where r is the corrected density of insertions in the 10-kb region centered on the ORF on the chromosome. For example, to assure that the probability P_0 that no transposon insertion is detected in the given gene by chance alone is <0.5 , we need the following: $rL > \log(2) = 0.639$. In our case, 604 of the 620 genes asserted to be essential satisfy this condition with rL of >0.639 , indicating that $\sim 97\%$ of all essential genes have a reliability of essentiality calls expressed by a P_0 of <0.5 . The number of essential genes with P_0 smaller than a fixed value is given in Table 1. A detailed list for each gene is presented in the supplementary data (see Table S1).

Identification of putative orthologs of *E. coli* genes in diverse set of microbial

genomes. Putative orthologs of *E. coli* genes were identified by using the ERGO database (<http://ergo.integratedgenomics.com/ERGO/>) (26). Protein families in ERGO correspond to homologous ORFs with identical assigned functions (24). With each update of the database, grouping of proteins into families is refigured through a multistep process including (i) formation of a family core from proteins corresponding to several ORFs that are bidirectional best FASTA hits for one another in their respective genomes, (ii) family extension by adding proteins with identical assigned functions and by performing FASTA searches (27) and adding matches with expectation values of less than a preset threshold, as described earlier (12), and (iii) refinement of a family grouping based on multiple ClustalW alignments (36) of all included sequences. To identify putative orthologs of *E. coli* proteins, all protein families in ERGO were automatically queried for the simultaneous presence of a protein(s) corresponding to an *E. coli* ORF(s) and proteins corresponding to ORFs from the genomes of 32 diverse bacterial species (*Agrobacterium tumefaciens*, *Anabaena* sp., *Aquifex aeolicus*, *Bacillus subtilis*, *Borrelia burgdorferi*, *Brucella melitensis*, *Buchnera* sp., *Campylobacter jejuni*, *Caulobacter crescentus*, *Chlamydia trachomatis*, *Clostridium acetobutylicum*, *Corynebacterium glutamicum*, *Deinococcus radiodurans*, *Fusobacterium nucleatum*, *Haemophilus influenzae*, *Helicobacter pylori*, *Listeria monocytogenes*, *Mesorhizobium loti*, *Mycobacterium tuberculosis*, *Mycoplasma pneumoniae*, *Neisseria gonorrhoeae*, *Pseudomonas aeruginosa*, *Ralstonia solanacearum*, *Rickettsia prowazekii*, *Sinorhizobium meliloti*, *Staphylococcus aureus*, *Streptococcus pneumoniae*, *Synechocystis* sp., *Thermotoga maritima*, *Treponema pallidum*, *Vibrio cholerae*, and *Xylella fastidiosa*). Results of this search were further supplemented by addition of ORFs from each of these genomes that are bidirectional best FASTA hits with corresponding *E. coli* genes.

Densities of essential genes and evolutionary retention indexes (ERIs) along the chromosome. The densities of essential genes along the *E. coli* chromosome (see Fig. 1B) were calculated within overlapping 100-kb regions displaced 1 kb from one another. For each 100-kb region, the essentiality was defined as the ratio of the number of essential genes to the total number of genes found in the region (N_E/N_T). The significance of essentiality for each 100-kb region was determined based on the hypergeometric distribution. Given that 620 of 4,291 *E. coli* genes were found to be essential, the probability of having N_E essential genes out of a total number of N_T genes within a 100-kb region is given by

$$P = \frac{\binom{620}{N_E} \binom{3,671}{N_T - N_E}}{\binom{4,291}{N_T}}$$

where $\binom{a}{b}$ denotes the number of ways to choose b out of a elements.

We determined the ERI for each of the 4,291 *E. coli* ORFs by calculating the fraction of genomes in the group that have an ortholog of the given ORF, with the number of representative organisms (N_G) equal to 33. Thus, if the number of organisms that contain an ortholog of the *E. coli* ORF is N_C , the ERI is given by the following formula: $\text{ERI} = N_C/N_G$. The ERIs along the *E. coli* chromosome were calculated within overlapping 100-kb chromosomal regions, displaced 1 kb from one another (see Fig. 1C). The ERI of each 100-kb region was determined by calculating the average of the ERIs for all ORFs located completely inside the region.

Data analysis within the context of system level metabolic organization. Using the information about the *E. coli* enzymes for all metabolic reactions available in the ERGO database, together with the essentiality data for the corresponding genes, we analyzed the correlation of enzyme essentialities within the known hierarchical structure of the *E. coli* metabolic organization. We have previously established a global topologic representation of the *E. coli* metabolic network, in which each branch on the hierarchical tree corresponds to a group of metabolites that are at its endpoints. Thus, each junction represents the module made up of the substrates that were clustered together up to that stage (28). For each branch, we can define an essentiality ratio based on the metabolic reactions present among the group of metabolites it represents.

To treat each reaction equally, we considered all links present between any two metabolites in the group, and for each of these links we took into account all the reactions that created the link. Specifically, for all pairs in the group, we included those metabolic reactions that transformed one of the substrates into another, according to a reaction list in which generic donor and acceptor moieties, such as H_2O and ATP, are not considered (see reference 28 for details) and to which an unambiguous insertion phenotype has been assigned (NR_{in}). Next, we counted those reactions whose corresponding catalytic enzymes proved to be essential ($\text{NR}_{\text{essential}}$). Note that since the hierarchical tree is constructed according to a two-step network complexity reduction procedure (28), there can be arcs between pairs of substrates that the tally does not include. To account for these, we

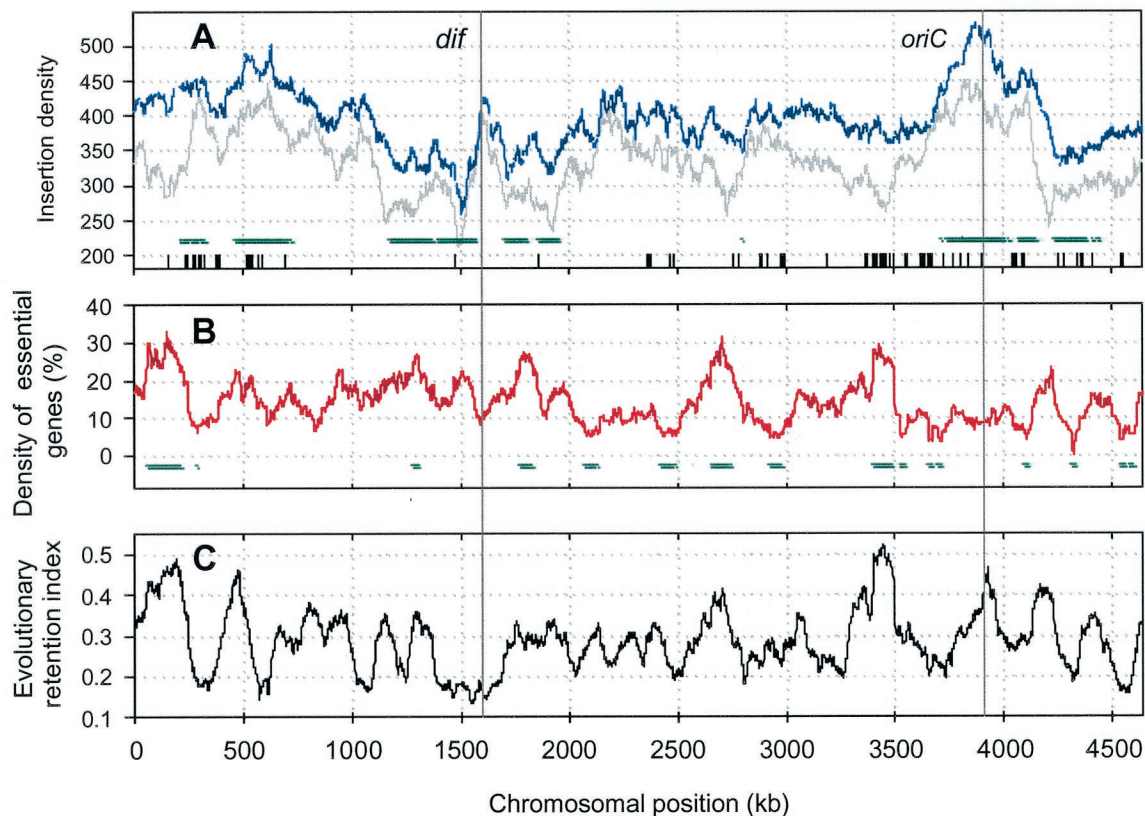


FIG. 1. Distribution of transposon insertion densities, densities of essential genes, and ERIs along the *E. coli* chromosome. (A) Gray lines show the transposon insertion densities calculated as the number of transposition events per 100-kb sliding window over the entire *E. coli* MG1655 chromosome. Values indicated by the blue lines were computed in a similar manner, except that all chromosomal regions corresponding to essential and ambiguous genes were excluded from the calculations in order to reconstruct insert distribution prior to selective outgrowth (see also Materials and Methods). Gaps in the data (chromosomal regions where transposition events could not be detected due to technical reasons) are indicated by short vertical lines along the *x* axis. These regions were excluded from all analyses. Nucleotide positions of the *E. coli* genome sequence correspond to those in reference 4. The regions where the distributions of transposition events significantly deviate ($P < 0.01$) from a Poisson process are marked by horizontal green lines. *oriC* shows the origin of chromosomal replication, and *dif* denotes the *dif* locus within the replication termination area. (B) Distribution of essential genes along the *E. coli* chromosome, defined as a percentage of essential genes in the total number of genes within a 100-kb-long chromosomal region (calculated per sliding window as described above). The regions where the numbers of essential genes significantly deviate ($P < 0.01$) from values that could arise by chance are marked by horizontal green lines. (C) ERIs along the *E. coli* chromosome, defined as the average ERI for all genes within each 100-kb region. The ERI for a gene is defined as the fraction of organisms in a diverse set of 33 bacterial species which contain an ortholog of the gene in their genomes.

examined each metabolic reaction with a known catalytic enzyme insertion phenotype on these internal arcs and incorporated them into the analysis. The essentiality of the branch (or module) is given by the fraction $NR_{\text{lethal}}/NR_{\text{all}}$ and represents the fraction of essential enzymes of all biochemical reactions within a given metabolic module (branch). For additional details, see the supplementary data.

TABLE 1. Number of essential genes with P_0 smaller than a fixed value

P_0	No. of essential genes	% of essential genes
<0.01	159	26
<0.05	281	45
<0.1	367	59
<0.2	476	77
<0.3	550	89
<0.4	587	95
<0.5	604	97
<0.6	610	98

RESULTS

Genome-scale genetic footprinting in *E. coli*. Genetic footprinting was first introduced for analysis of gene essentiality in *S. cerevisiae* (33). A modification of this technique using a Tn5-based in vitro transposome system (11) in *E. coli* was previously described, and gene essentiality within three cofactor biosynthetic pathways has been analyzed (9). Here we have extended this pilot analysis to the whole-genome level by using the same standardized growth conditions. The general experimental scheme is illustrated in the supplementary data. Briefly, following transposon mutagenesis, a population of $\sim 2 \times 10^5$ independent mutants was grown aerobically for 23 doublings in enriched LB medium supplemented with kanamycin. Genomic DNA was isolated from the whole population and used to map individual transposon inserts with a nested PCR approach.

Distribution of the 1.8×10^4 distinct insert locations detected along the *E. coli* chromosome is illustrated in Fig. 1A.

TABLE 2. Distribution of essential and nonessential genes and average ERIs in selected functional categories^a

Functional category	Description	Total no. of ORFs	No. E	No. N	No. ?	No. ND	% E	Mean ERI
AAM	Amino acid metabolism	138	21	108	1	8	15	0.5
CHM	Carbohydrate metabolism	219	21	178	3	17	10	0.4
NCM	Nucleotide and cofactor metabolism	181	53	119	2	7	29	0.5
LPC	Lipid, lipopolysaccharide, lipoprotein, peptidoglycan, and cell wall biosynthesis	126	34	73	5	14	27	0.5
NAM	Nucleic acid metabolism	156	43	96	5	12	28	0.6
PMS	Protein metabolism and secretion	167	80	57	18	12	48	0.7
MSM	Miscellaneous metabolism	94	14	72	2	6	15	0.4
BEN	Bioenergetics	100	15	75	4	6	15	0.3
SMC	Signaling, motility, and chemotaxis	153	12	125	3	13	8	0.3
RCD	Expression regulation and cell cycle and division	177	30	118	13	16	17	0.2
MTR	Membrane transport	276	22	244	3	7	8	0.3
PHT	Phage- and transposase-related processes	62	12	35	2	13	19	0.3
CAT	All categorized	1,849	357	1,300	61	131	19	0.4
UNC	Uncategorized	2,442	263	1,826	157	196	11	0.2
Total	Categorized and uncategorized	4,291	620	3,126	218	327	14	0.3

^a Abbreviations are as follows: E, essential; N, nonessential; ND, not determined; ?, ambiguous.

The densities of transposon insertion events are randomly distributed, with two notable exceptions: an overall maximum around the origin of replication (*oriC*) and a minimum around the terminus (*dif*). This may reflect increased target copy number at the origin of replication in the actively dividing bacterial population used in this experiment. The overall insertion density is 3.218/kb, without appreciable variation between coding (3.221/kb) and noncoding (3.193/kb) regions.

Assessment of conditional gene essentiality based on genetic footprinting data. Unambiguous essentiality assessments were made for 3,746 (or 87% of the total) *E. coli* protein-encoding genes or ORFs (Table 2). Of these, 620 (14%) were asserted to be essential, and 3,126 (73%) were asserted to be nonessential (dispensable) based on the occurrence of transposon inserts within each ORF and the overall insertion density in the local environment, as described in the supplementary data. The complete essentiality list is reported in the supplementary data (see Table S1). No assertions could be made for 327 genes for technical reasons, such as limited efficiency of PCRs in certain regions of the *E. coli* chromosome or nonspecific primer annealing in areas of DNA repeats. For 218 genes, we considered the evidence to be insufficient for a specific conclusion about essentiality. These genes were systematically called ambiguous, according to the criteria listed in the supplementary data. For example, ORFs shorter than 240 bp (<80 aa) and with no inserts were consistently classified as ambiguous rather than essential. In certain cases, relatively long ORFs (>900 bp) containing only a single transposon were designated ambiguous rather than nonessential.

Our results are generally consistent with previously published data on individual genes and with data from currently available collections of systematic gene deletions in *E. coli*. For example, of the 1,379 individual gene deletion mutants listed at the University of Wisconsin *E. coli* Genome Project website (<http://www.genome.wisc.edu/functional/tmmutagenesis.htm>), only 12% produced apparently conflicting designations of genes as essential (for a detailed list of the discrepancies, see

Table S2 in the supplementary data). Although we have not attempted to reconcile each individual case, several reasons for discrepancies can be envisioned. Most importantly, the term essential, which intuitively suggests an absolute requirement for cell viability, also applies to any gene that imparts a substantial fitness advantage. Thus, mutants lacking gene products necessary for maintaining vigorous growth fall into the same category as those with “true lethal” mutations. Therefore, certain genes may be classified as essential by genetic footprinting, yet corresponding viable deletion mutants may be obtained. In addition, differences in medium compositions, aeration levels, temperatures, and cell densities may account for many inconsistencies. Surprisingly, polar effects, in which transposon insertion into dispensable genes disrupts transcription of essential genes, are relatively rare in genetic footprinting. This may be due to the presence of weakly active promoter-like sequences within the transposon used in these experiments (9, 11). Most examples of polar effects are associated with genes that may require high levels of expression to sustain rapid growth rates.

Discrepancies resulting from inserts detected in the genes otherwise considered to be essential also occur. In some cases, single inserts occur close to protein termini or in interdomain boundary regions in multidomain proteins. For proteins consisting of two or more independently functioning domains, inserts may be tolerated within the 3' portion of the gene if the C-terminal domain of the protein it encodes is associated with a dispensable function. This can occur even when a function associated with the N-terminal domain (from the 5' region of the gene) is genuinely essential (as with *ftsX* [9]). Small, localized chromosomal duplications may account for inserts in genes otherwise recognized as essential (2). In this scenario, one copy of a duplicated gene provides the essential function while the other copy containing the transposon is stabilized by selection for kanamycin resistance. Large genes with only a small number of inserts may fall into this category since the

total number of specific duplications within the population prior to transformation is probably very small (25).

Functional context analyses of essentiality data. The interpretation of genomic essentiality data can be approached in a number of alternate ways, such as by using chromosomal (positional), functional (system level), or phylogenetic (evolutionary) context analysis. In addition to refining initial essentiality assignments and reconciling apparent discrepancies with existing knowledge, such analyses can improve and expand existing understanding of the systemic behavior of the cell at various levels. Without attempting a comprehensive analysis, we have limited the scope of our efforts to (i) prototyping and illustrating such analysis by using selected examples from various functional systems, (ii) evaluating the internal consistency of our data, and (iii) developing preliminary observations at the system level, as presented below.

Initially, we analyzed the data in a functional context, which involved dividing the overall physiology of the organism into smaller, internally coherent subsystems such as amino acid biosynthesis, nucleotide metabolism, and other broad functional categories (Table 2). This approach mirrors the standard didactic subdivision of microbial biochemistry and physiology. It also provides an organizational framework with which to analyze total genomic data and allows specific metabolic questions to be addressed.

For consistency, our functional analysis is based exclusively on SWISS-PROT functional annotations (8). Each of the 1,849 gene products with specific SWISS-PROT annotations and defined biochemical functions supported by solid experimental evidence was placed into one of the 12 functional categories (Table 2 and supplementary data [see Table S1]). Among the remaining 2,242 uncategorized protein-encoding genes, many have been tentatively annotated in SWISS-PROT and other databases, but most of these annotations either fall short of giving a specific testable function or have not been confirmed by direct experiments. As expected, the ratios of essential genes within various functional categories are rather uneven (Table 2). Categories that include gene products involved with key aspects of cellular metabolism (such as nucleic acid and protein metabolism) contain a substantially higher percentage of essential genes (28 and 48%, respectively) than the average for the entire genome (14%). The percentages of essential genes in categories such as signaling, motility, and chemotaxis (8%) and membrane transport (8%) are substantially below the whole-genome benchmark. The average essentiality for the subset of 2,242 uncategorized genes (11%) is substantially lower than the average for the subset of categorized genes (19%). Several representative metabolic and nonmetabolic systems (7 of 12 functional categories) were selected for use as examples of functional context analysis and for evaluation of the internal consistency of the data. Here we describe one such analysis, with additional detailed interpretations presented in the supplementary data.

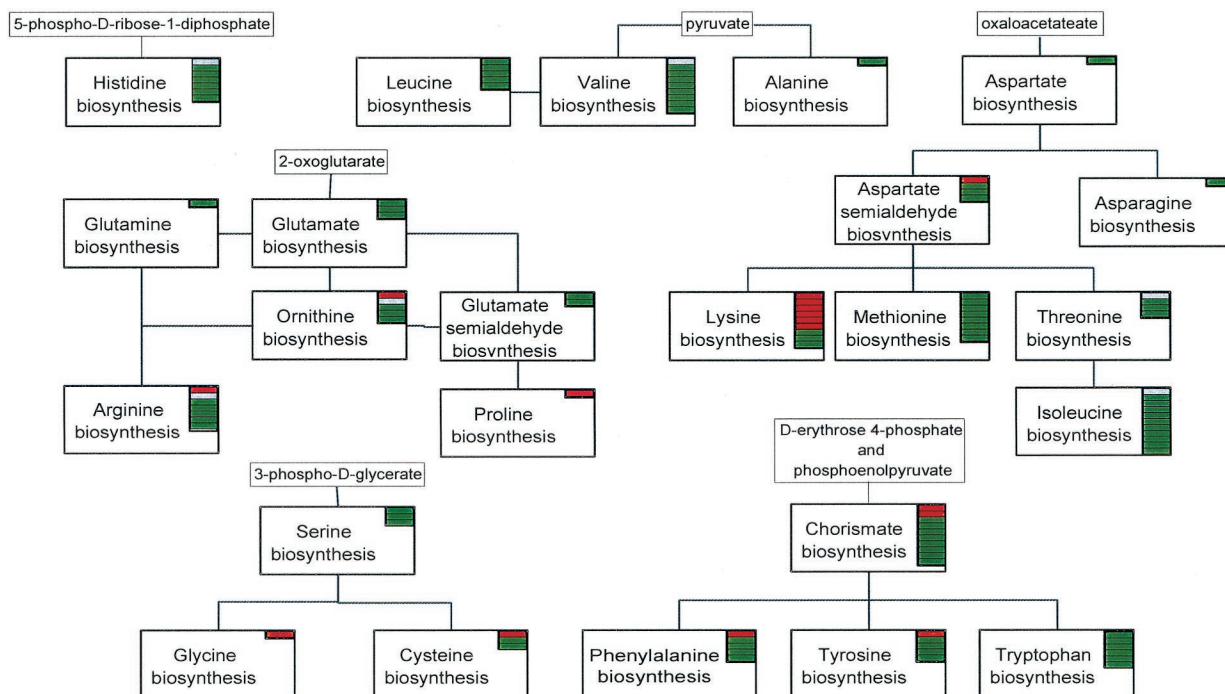
Amino acid metabolism: lysine biosynthesis. Most of the genes responsible for biosynthesis of various amino acids were expected to be nonessential since the medium contains most of the amino acids required for growth. With a few notable exceptions, this expectation was confirmed by our results. Of the 91 genes with specific SWISS-PROT annotations indicating involvement in amino acid biosynthesis, only 16 appear to be

essential (Fig. 2A). Six of these genes are involved in lysine biosynthesis. *E. coli* produces lysine from aspartate via the nine-step pathway (Fig. 2B). Although lysine is available in the growth medium, its immediate precursor, diaminopimelate (DAP), which is required for cell wall biosynthesis, is not. The *lysA* gene encoding the enzyme that converts DAP to lysine at the last step of this pathway is dispensable. Analysis of DAP-lysine biosynthesis provides an example of refining pathway reconstruction and individual functional assignments based on genome-scale essentiality data. Genes (*asd*, *dapA*, *dapB*, *dapD*, *dapE*, and *dapF*) encoding most of the enzymes leading to DAP production are essential. The first gene in this pathway (*lysC*), encoding aspartokinase III, is dispensable due to the functional redundancy of the additional aspartokinase isozymes (encoded by *metL* and *thrA*). In contrast, the *asd* and *dapA* genes involved with the second and the third steps of DAP-lysine biosynthesis are essential in spite of the existence of apparent paralogs. Proteins encoded by the *yjhH* and *yagE* functionally uncharacterized genes are often annotated as potential dihydrodipicolinate synthases based on their high sequence similarities with the *dapA* gene product (BLAST E scores of $4e^{-33}$ and $2e^{-28}$, respectively). However, genetic footprinting data suggest that under our experimental conditions neither is capable of complementing loss of the essential *dapA* function. The opposite situation is observed with succinyl-DAP aminotransferase (encoded by *argD*), which is firmly defined as dispensable in our data. This apparent inconsistency can be resolved by assuming functional complementation by the *argM* gene product. The *argM* gene is known to encode succinyl-ornithine transaminase, which is primarily involved in arginine biosynthesis. However, this enzyme is closely related to succinyl-DAP aminotransferase by sequence, and the aminotransferases are known to possess rather broad substrate specificities, especially for structurally similar substrates (such as succinyl-DAP and succinyl-ornithine). Overexpression of the *argM* gene has been demonstrated to suppress an *argD* mutation in *E. coli* (32).

Phylogenetic analysis of essentiality data within functional groups. To assess the data set from an evolutionary perspective, we examined the distribution of conditionally essential and dispensable *E. coli* genes with respect to the occurrence of putative orthologs across a broad range of diverse bacterial genomes. Putative orthologs within a reference set of 32 complete bacterial genomes chosen to represent maximum phylogenetic diversity were identified based on protein families, supplemented by bidirectional best hits (see Materials and Methods). For this analysis we introduce a simple parameter: an ERI computed for each *E. coli* gene as the fraction of genomes from the reference set containing a putative ortholog of the gene. ERI values varying from 0 (for genes unique to *E. coli*) to 1.0 (for omnipresent genes) are provided in the supplementary data (see Table S1). In a recent study, the Profiling of *E. coli* Chromosome data (<http://www.shigen.nig.ac.jp/ecoli/pec>) were used to demonstrate a remarkable tendency of essential gene sequences to be more evolutionarily conserved than those of nonessential genes (19). In our analysis, we used ERI values to focus on occurrence of essential and nonessential genes (preservation of orthologs) rather than on conservation of their respective sequences.

Figure 3A depicts the overall number of *E. coli* genes in

A



B

Reaction	Step #	Enzyme	Gene name	Essentiality
Aspartate				
↓	1	Aspartate kinase	lysC thrA metL	Non-Essential Non-Essential Non-Essential
Phosphoaspartate				
↓	2	Aspartate-semialdehyde dehydrogenase	asd usg	Essential Non-Essential
Aspartate semialdehyde				
↓	3	Dihydrodipicolinate synthase	dapA yagE yjhH	Essential Non-Essential Non-Essential
Dihydrodipicolinate				
↓	4	Dihydrodipicolinate reductase	dapB	Essential
Tetrahydrodipicolinate				
↓	5	2,3,4,5-tetrahydropyridine-2-carboxylate n-succinyltransferase	dapD	Essential
Succinyl-L-2-amino-6-keto-pimelate				
↓	6	Succinyldiaminopimelate aminotransferase	argD argM	Non-Essential Non-Essential
Succinyl-L-2,6-diaminopimelate				
↓	7	Succinyl-diaminopimelate desuccinylase	dapE	Essential
L,L-2,6-diaminopimelate				
↓	8	Diaminopimelate epimerase	dapF	Essential
Meso-2,6-diaminopimelate				
↓	9	Diaminopimelate decarboxylase	lysA	Non-Essential
Lysine				

FIG. 2. Essentiality of genes controlling amino acid biosynthesis in *E. coli*. (A) Functional overview of amino acid biosynthesis. Each block represents one or more pathways leading to production of a particular amino acid or its key intermediates (shown in smaller boxes). Within each block, stacked bars represent the gene products involved in the pathway (according to SWISS-PROT release of June 2002). Bars are colored according to gene essentiality (green, nonessential; red, essential; gray, undefined). (B) Detailed representation of the lysine biosynthetic pathway. Genes predicted in the ERGO database to be paralogs in this pathway are shown, in addition to genes whose roles in the biosynthesis of lysine have been experimentally verified (in bold).

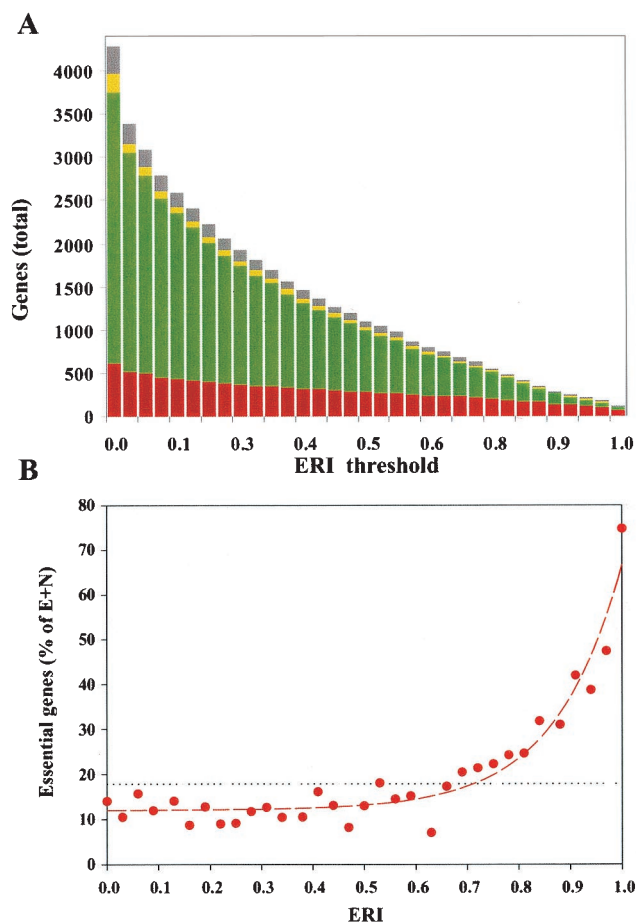


FIG. 3. Distribution of *E. coli* genes as a function of ERIs. (A) Total number of genes with an ERI above the threshold plotted versus the ERI threshold. Color coding within bars represents fractions of essential (red), nonessential (green), ambiguous (yellow), and missing (gray) genes for each incremental increase of ERI threshold (with 33 diverse genomes in the reference set). (B) Fractions of essential genes at different ERI values. The data were fitted with the following function: $y = y_0 + ae^{bx}$, where y_0 is 12.0 ± 0.9 , a is 0.023 ± 0.019 , and b is 7.8 ± 0.8 (dashed red line). The dotted line represents the fractions of essential genes for the whole genome. (The fractions plotted are defined as the number of essential genes versus the number of essential (E) and nonessential (N) genes. Unknown or ambiguous genes are not taken into account.)

decreasing order over the range of ERI values. An initial sharp decrease in the number of preserved genes ($\sim 40\%$) occurs over a rather small phylogenetic distance of less than four genomes in our reference set ($\text{ERI} \leq 0.1$). Further decay is at much lower rates, and orthologs of $\sim 10\%$ of *E. coli* genes are preserved in at least 25 diverse genomes ($\text{ERI} \geq 0.8$). This reflects a nonrandom ortholog preservation pattern, characterized by a highly conserved core group of genes. This core is highly enriched by genes identified as essential in our study. The tendency of essential genes to be evolutionarily preserved is also reflected in Fig. 1, demonstrating a significantly positive correlation (0.5240) between essentiality (Fig. 1B) and ERIs (Fig. 1C) along the *E. coli* chromosome. Similarly, plotting the fraction of essential genes at different ERI values demonstrates that the relationship between the two parameters has

the following form: $y = y_0 + ae^{bx}$, implying that the essentiality of genes with a given ERI is due partly to a very strong tendency of essential genes to be retained by evolution (the exponential behavior dominant above an ERI of 0.6) and partly to an essential gene fraction of $\sim 10\%$ that is present among genes within any ERI value group (Fig. 3B).

Comparison of average essentiality and ERI values between different functional categories reveals significant correlation (Table 2). Functional categories including highly specialized proteins such as transporters, regulators, and signaling molecules are characterized by average ERI values close to the average for the whole genome (~ 0.3). Average essentiality within these groups also does not exceed an overall whole-genome level ($\sim 14\%$). The least essential group of all uncategorized proteins with historically elusive functions has the lowest average ERI, ~ 0.2 . Therefore, many of these proteins are likely to be specific to the environmental and phylogenetic niches of *E. coli*. On the other hand, the bulk of cellular intermediary metabolism (categories AAM, CHM, NCM, LPC, and MSM [Table 2]) is associated with ERI values of 0.4 to 0.5. Essentiality within these metabolic categories varies depending on the levels of functional redundancy of their constituents in rich medium. Not surprisingly, the highest ERI values (up to 0.7) as well as the highest ratio of essential genes (up to 48%) occurs in functional categories that include replication, transcription, and translation, i.e., cellular processes that are conserved and unconditionally essential in most organisms.

Figure 4 illustrates the changes in distribution of essential genes between functional categories depending on their tendencies to be evolutionarily preserved. An initial bias in distribution of all categorized essential genes towards those involved with synthesis and processing of informational macromolecules increases dramatically at higher ERI values. The fraction of all essential genes contributed jointly by the functional categories PMS and NAM (Table 2) ($\sim 30\%$) increases almost twofold (up to $\sim 60\%$) for a subset of essential genes with ERIs of >0.8 , ultimately exceeding 90% as the ERI approaches 1.0.

This analysis reveals two distinct classes of essential genes, which may be referred to as broadly preserved essential genes and species-specific essential genes. A subset of less than 180 genes ($\sim 4\%$ of the genome) with ERIs of >0.8 accounts for $\sim 25\%$ of all of the essential genes revealed in this study, and it appears to provide an approximation of broadly preserved essential genes. Functional content analysis of this subset (Fig. 5) strongly supports the expectation that these genes represent universally and unconditionally essential constituents of cellular central machinery. This notion is in good agreement with available complete and partial gene essentiality datasets for *Mycoplasma pneumoniae* and *Mycoplasma genitalium* (15), *Haemophilus influenzae* (1), *Staphylococcus aureus* (7, 18), and *Streptococcus pneumoniae* (35). The overwhelming majority (70 to 87%) of assigned genes in these data, which correspond to *E. coli* genes listed in Fig. 5, appear to be essential (see Table S5 in the supplementary data for details). Of note, many of these broadly preserved essential genes, including those with yet undefined functions, may be considered potential broad-spectrum anti-infective drug targets (9, 29).

In contrast, more than 75% of genes within the set of spe-

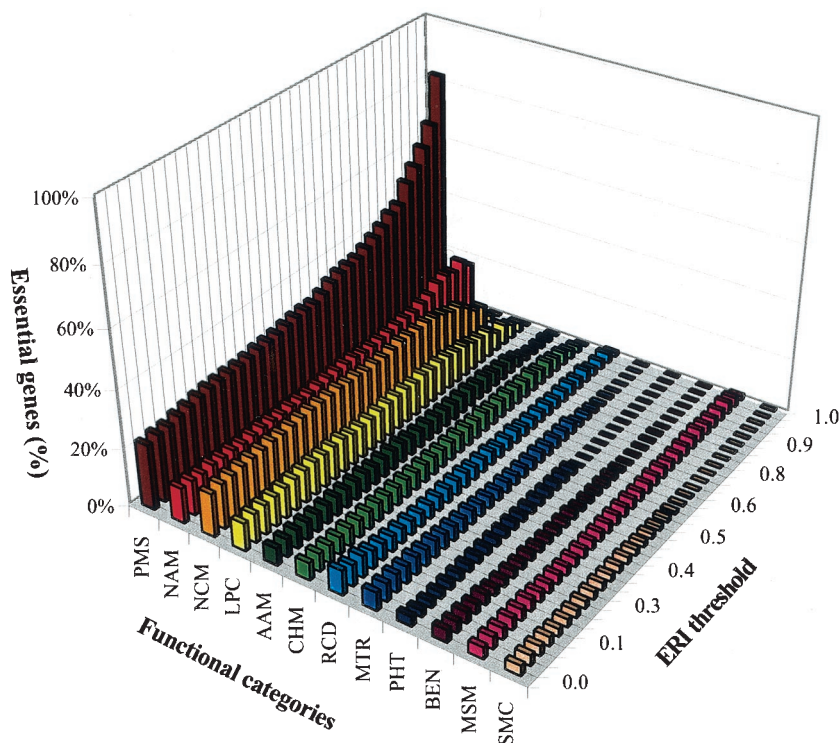


FIG. 4. Distribution of essential genes among functional categories as a function of ERI thresholds. Functional categories are color coded and specified by three-letter designations as in Table 2. Within every threshold group, each bar represents the fraction (percent plotted on y axis) of all categorized essential genes corresponding to the number of essential genes in a given category (x axis) with ERI values above the set threshold (z axis).

cies-specific essential genes (which account for $\sim 30\%$ of all essential *E. coli* genes with ERI values of <0.1) encode uncategorized proteins with poorly defined or completely unknown functions. Many of the genes with known functions within this class are related to transcription regulation, membrane transport, signaling, and other cellular processes whose essentiality is either strictly condition dependent or limited to a set of very specific needs of *E. coli* and closely related species.

Among the 263 essential genes marked in our analysis as uncategorized (see Table S1 in the supplementary data), 19 genes have specific functions assigned to them while 73 genes have putative assignments (according to SWISS-PROT and other public archives). Those include assignments indicating just an element of possible function, such as “probable GTP-binding protein” (*ychF*). For the remaining 171 genes, we were unable to find any reliable functional assignments. These genes may be qualified as essential unknowns (at least at the time when this analysis was performed). The list of these genes along with their respective ERI values is provided in the supplementary data (see Table S6). Only 10 (*yjiL*, *yjeE*, *ybeY*, *yebC*, *yjgF*, *ydeE*, *yoaB*, *yqgF*, *ycdK*, and *yhbC*) of the essential unknowns ($<6\%$) are broadly conserved in bacteria (ERIs of 0.8 to 1). In contrast, more than 60% of genes in this set are poorly conserved across our reference set of diverse genomes (108 genes with ERIs of 0 to 0.1). Less than half of them (42 genes) are conserved in most *Enterobacteriaceae*, while others are present only in *E. coli* and some closely related species.

System level analysis of essentiality data within topologic modules of *E. coli* metabolism. It is widely recognized that the thousands of components of a living cell are dynamically interconnected, so that cellular functional properties are a result of the complex intracellular web of molecular interactions within the cell (14, 22, 23). This is perhaps most evident with intermediary metabolism, in which hundreds of metabolic substrates are densely integrated through biochemical reactions (17). Metabolic networks are organized into many small, highly connected topologic modules that combine in a hierarchical manner into larger, less cohesive units, with their numbers and degrees of clustering following a power law, as previously demonstrated for 43 reference organisms (28). Within *E. coli*, hierarchical modularity closely overlaps with known metabolic functions (28).

To comprehend the results of individual gene essentiality in the context of cellular system level functional organization, we projected the essentiality phenotype of metabolic enzymes onto a global topologic representation of the *E. coli* metabolic network (28). As shown in Fig. 6, the overall essentiality ratio of metabolic enzymes within the full metabolic network is relatively low, with essential enzymes limited to a subset of modules. Visual inspection of the figure indicates that while many metabolic modules are almost entirely nonessential, at the lowest hierarchical level several branches corresponding to small topologic modules appear to be essential, i.e., they are composed of biochemical re-

94 INFORMATION			58 METABOLISM		
69 PMS			21 NCM		
16	Amino acyl tRNA synthesis	<i>alaS argS aspS</i> <i>cysS gltX hisS</i> <i>ileS leuS metG</i> <i>pheS proS serS</i> <i>thrS trpS tyrS</i> <i>valS</i>	9	NTP biosynthesis	<i>pyrG pyrH adk</i> <i>nrdA cmk thyA</i> <i>gmk tmk ndk</i>
29	Ribosome	<i>frr rplB rplC</i> <i>rplD rplF rplJ</i> <i>rplL rplM rplN</i> <i>rplO rplR rplS</i> <i>rplT rplW rplX</i> <i>rplY rpsB rpsE</i> <i>rpsF rpsH rpsI</i> <i>rpsJ rpsL rpsM</i> <i>rpsN rpsO rpsP</i> <i>rpsQ rpsS</i>	5	FMN/FAD biosynthesis	<i>ribH ribE ribD</i> <i>ribA ribF</i>
			3	CoA biosynthesis	<i>dfp coaD coaE</i>
			2	NAD/NADP biosynthesis	<i>nadE ppnK</i>
			1	Folate biosynthesis	<i>folK</i>
			1	Other cofactors	<i>metK hemC trx</i>
			20 LPC		
			4	Fatty acid biosynthesis	<i>fabH fabG fabD</i> <i>birA</i>
			6	Isoprenoid biosynthesis	<i>ispB ispE dxs</i> <i>ispA ispG uppS</i>
			5	Peptidoglycan biosynthesis	<i>ftsI ddlB murC</i> <i>murE murG</i>
12	Translation	<i>def efp fnt</i> <i>fusA infB infC</i> <i>map pth tsf</i> <i>tufA tufB hemK</i>	3	Phospholipid biosynthesis	<i>pgsA cdsA plsC</i>
			2	Lipoprotein biosynthesis	<i>lolD lgt</i>
8	Sec-dependent secretion	<i>ffh ftsY grpE</i> <i>lepB lspA secA</i> <i>secD secY</i>	9 CHM		
			6	Glycolysis/TCA	<i>pgk zwf gapA</i> <i>tktA ack lpdA</i>
4	Protein folding	<i>dnaK groL groS</i> <i>ppiB</i>	2	Aminosugar biosynthesis	<i>glmS glmU</i>
			1	PRPP Biosynthesis	<i>prsA</i>
19 NAM			8 AAM		
9	Replication	<i>dnaA dnaB dnaE</i> <i>dnaG dnaX gyrA</i> <i>priA rnhB topA</i>	3	DAP biosynthesis	<i>asd dapA dapB</i>
			1	D-Glu biosynthesis	<i>murI</i>
1	Repair	<i>ligA</i>	1	Se-Cys biosynthesis	<i>iscS</i>
6	RNA processing/modification	<i>rimM znc rluD</i> <i>miaA trmU trmD</i>	3	Other amino acids	<i>aroK glyA proC</i>
			19 UNCHARACTERIZED		
3	Transcription	<i>pnp rpoB rpoC</i>	7	GTP/ATP hydrolase-related	<i>engA era trmE</i> <i>yebC ychF yhbZ</i> <i>yjeE</i> <i>ybeY yqgF yeaZ</i>
6 RCD			3	Hydrolase-related	<i>yhbF</i>
4	Cell division	<i>ftsA ftsW ftsZ</i> <i>mraW</i>	1	Methyltransferase-related	<i>yoaB yjgF ycdK</i>
			3	Translation-related	<i>msbA mviN ydeE</i>
2	Transcription factors	<i>nusB xpoD</i>	5	Membrane proteins	<i>yhbG yjeE</i>
			1	RNA modification-related	<i>yciL</i>

FIG. 5. *E. coli* genes found to be essential and preserved in over 80% of diverse bacterial genomes (ERI > 0.8). These universal essential genes are grouped by functional categories (described in Table 2). NTP, nucleotide triphosphate; FMN, flavin mononucleotide; FAD, flavin adenine dinucleotide; CoA, coenzyme A; TCA, tricarboxylic acid cycle; PRPP, phosphoribosyl pyrophosphate.

actions catalyzed by predominantly essential enzymes. Of these, the largest fractions are within the topologic modules related to nucleotide, coenzyme, and lipid metabolism. The pyrimidine metabolic module appears to contain the highest level of essential reactions.

A significant correlation between essentiality and ERI values is apparent within metabolic modules, and many of the highly essential modules also contain metabolic enzymes with the highest ERI values (Fig. 6). Generally, essentiality and evolutionary retention of metabolic enzymes correlate, although exceptions are also evident as illustrated in detail for the pyrimidine module (supplementary data [see Fig. S3]). Pyrimidine metabolism, however, represents a special case in *E. coli* MG1655, since the *rph-1* mutation in this

strain depresses expression of the downstream *pyrE* gene (16). This strain is prototrophic for pyrimidines but grows significantly better in uracil-supplemented media. Although our studies were performed with rich media containing significant amounts of exogenous pyrimidines, the low level of *pyrE* transcription may have affected the ability of cells to efficiently adjust the relative levels of the pyrimidine nucleotides. This may explain the relatively high level of gene essentiality within the pyrimidine-related topologic module. These observations, however, may also reflect a hypothesized generic feature of metabolic networks: their limited ability to fully compensate for perturbations by reorganization of metabolic fluxes within evolutionarily conserved topologic modules.

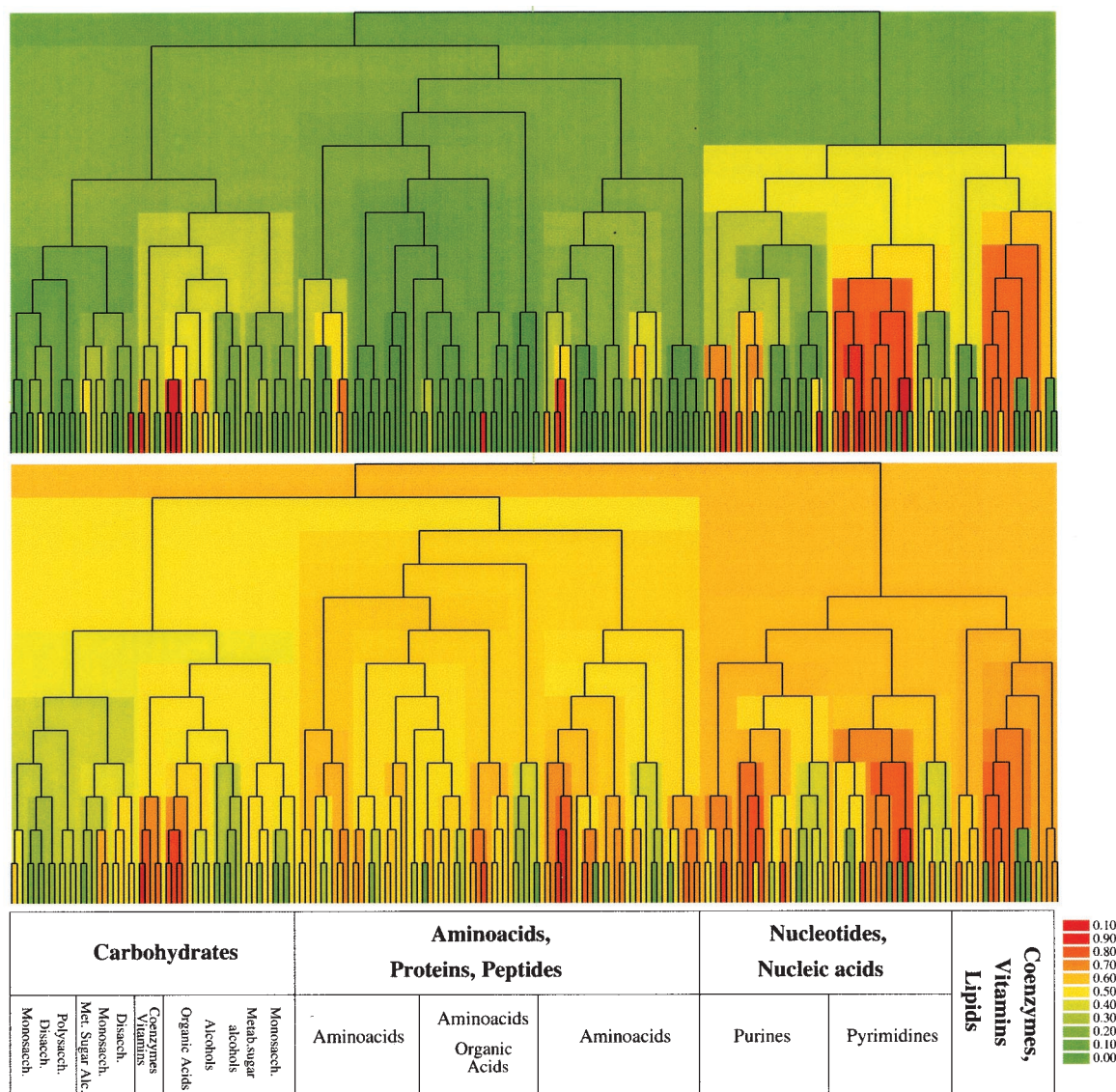


FIG. 6. The evolutionary retention and essentiality ratio of enzymes in the topologic modules of *E. coli* metabolism. The hierarchical tree derived from the topologic overlap matrix of *E. coli* metabolism that quantifies the relation between the various modules is shown, as previously described (28). The branches of the tree are color coded according to the fraction of essential enzymes (top panel) and the average ERI score of enzymes (bottom panel) catalyzing the biochemical reactions within a given topologic module. Red indicates a 100% essentiality/conservation ratio within a module. Note that essentiality is not uniformly distributed across all modules (branches), but we observe a few small modules with very high fractions of essential enzymes, while the majority of modules contain no or only a few essential enzymes. A similar segregation of modules with high evolutionary conservation is observed in the second panel, with their locations often correlating with those of the high essentiality modules. The predominant biochemical classes of substrates used to group the metabolites are shown. Polysacch., polysaccharide; disacch., disaccharide; monosacch., monosaccharide; met. sugar alc., metabolic sugar alcohols.

DISCUSSION

A genetic footprinting technique was used to assess gene essentiality in *E. coli* K-12 across the entire genome under uniform growth conditions (logarithmic aerobic growth of strain MG1655 in enriched LB medium). This approach generated an internally coherent data set, which was examined at increasingly abstract levels to refine models of cellular organization. At the finest level, individual gene essentiality reveals basic physiologic information about cellular metabolism under specific growth conditions. At a more abstract level, the data

can be used for focused comparative genomic analysis to define the core bacterial genetic repertoire, while at the highest level of abstraction, the data can be used to detect organizational principles of cellular networks.

Functional context analysis based on projection of the gene essentiality data across a whole-genome functional reconstruction (metabolic and nonmetabolic pathways and networks) provides a powerful way to refine and interpret the results of genetic footprinting. This type of analysis, previously described only for a limited set of metabolic pathways (9) and extended

here to the whole-genome level, reveals a remarkable consistency between experimental observations and our present understanding of biochemical pathways and individual gene functions. Based on the overall consistency, one can resolve ambiguities, reconcile conflicting essentiality data, and even make tentative assignments for individual uncharacterized genes if they occur within well-known functional contexts (pathways).

Additionally, functional context analysis improves and extends our understanding of the systemic behavior of the cell at all levels: from individual genes and gene products to large functional systems and networks. Global projection of experimentally determined gene essentiality over a functional reconstruction model bridges the gap between two fundamentally different but related concepts: essential functions and essential genes. For example, essentiality data can distinguish functional (mutually complementing) and nonfunctional (noncomplementing) paralogs of genes with essential functional roles.

Analysis of essentiality data in a physiological context as a function of various factors and conditions, such as medium composition, aeration, growth phase, and temperature, etc., provides an opportunity to connect large functional modules with particular types of physiological states. Performing such analyses for a variety of conditions will provide critical support to systemic modeling efforts, such as flux-balance (6) and elementary mode analyses (34), and to our understanding of topologic modules (28). In this respect, the unexpected number of essential enzymes within the pyrimidine metabolic module in a *pyrE*-challenged *E. coli* strain reveals a significantly reduced ability of this module to tolerate additional gene inactivation, even in rich media. This suggests that the capacity for reorganization of metabolic fluxes within evolutionarily conserved, and presumably universally important, metabolic modules may be reduced, as a consequence either of their less evolved connectivity (37) or the performance of their functions at near optimality with corresponding innate fragility to uncommon error (5). The validity of these hypotheses will need to be tested by future experiments.

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